

Anexa nr. 1  
La Procedurile administrative pentru notificarea  
dispozitivelor medicale care dețin marcajul CE  
Către Agenția Medicamentului  
și Dispozitivelor Medicale

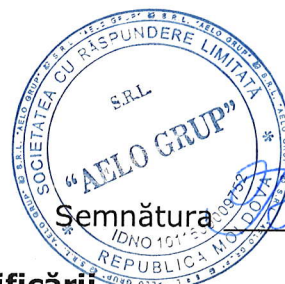
**NOTIFICARE**  
pentru înregistrarea dispozitivelor medicale în Registrul de stat  
al dispozitivelor medicale  
nr. .... din .....

Solicitantul **AELO GRUP SRL**, cu sediul **mun.Chișinău, str.Mitropolit Petru  
Movila 23/5, ap.6,**  
tel./fax: **+373 61 033 993/022 60 14 91**, e-mail: **aelogrup@gmail.com**,  
solicit înregistrarea în Registrul de stat al dispozitivelor medicale a următoarelor  
categorii și tipuri de dispozitive medicale pentru introducerea și punerea la dispoziție  
pe piață a:

**Defibrilator Monitor Comen S5**

Se anexează următoarele acte:  
CE;ISO;Decalaratie de conformitate;Brosura

Data 20.06.2023



Semnătura

**Tabelul de recepționare a notificării**

(se completează de către Agenție în momentul depunerii notificării de către  
solicitant)

Comentarii cu privire la acceptul/refuzul recepționării notificării, inclusiv motivul refuzului	
Data/nr. de ordine atribuit notificării de către Agenție (în cazul acceptării recepționării)	
Numele, prenumele, funcția persoanei responsabile de recepționarea dosarului	
Semnătura persoanei responsabile	

Către Agenția Medicamentului și Dispozitive Medicale

**DECLARAȚIE PE PROPRIE RĂSPUNDERE**

**Solicitant: Aelo Grup SRL,**

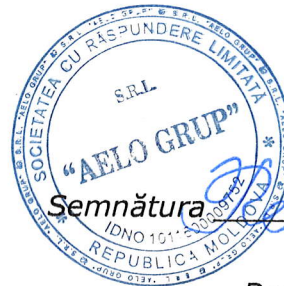
**cu sediul în: mun.Chișinău, str.Mitropolit Petru Movila 23/5, ap,6**

declar pe proprie răspundere, cunoscând prevederile art. **352<sup>1</sup>**, Codul Penal al Republicii Moldova cu privire la falsul în declarații, că documentele și datele furnizate pentru notificarea dispozitivului medical:

**Defibrilator Monitor Comen S5**

**Sunt autentice și corespund realității.**

*Cobzari -Țurcan Rodica*  
*Administrator*



*Data 20.06.2023*

## EC Declaration of Conformity

**Manufacturer:**

Shenzhen Comen Medical Instruments Co.,Ltd.

**Address:**

Floor 10, Floor 11 and Section C of Floor 12 of  
Building 1A & Floor 1 to Floor 5 of Building 2,  
FIYTA Timepiece Building, Nanhuan Avenue,  
Matian Sub-district, Guangming District,  
Shenzhen, Guangdong, 518106, P.R. China.

**Whose Single Authorized Representative:**

Lotus NL B.V.

**Address:**

Koningin Julianaplein 10, 1e Verd, 2595AA,  
The Hague, Netherlands.

SRN: NL-AR-000000121

We, the manufacturer(SRN number: CN-MF-000002236),, declare at our sole responsibility  
that following products

Product name	Model	Basic UDI-DI
Defibrillator Monitor	S8, S6, S5, S3	69454290DM001K6

meet the provisions of Directive 93/42/EEC

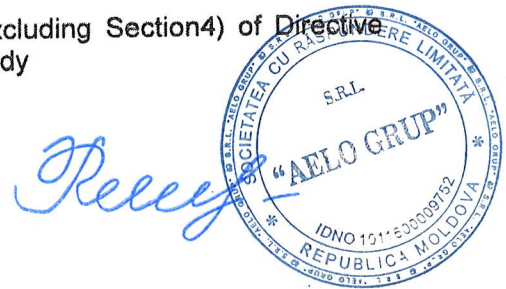
The medical device has been assigned to class IIb according to Rule 9 and Rule 10 in Annex  
VIII of Directive 93/42/EEC. It bears the mark

CE 1639

The product concerned has been designed and manufactured under a quality management  
system according to Annex II (excluding Section4) of Directive 93/42/EEC.

Compliance of the designated product with the Annex II (excluding Section4) of Directive  
93/42/EEC has been assessed and certified by the Notified Body

**SGS Belgium NV**  
**SGS House Noorderlaan**  
**87 2030 Antwerp Belgium**  
CertificateNo.: CN19/41057  
Issuedate: 2021.03.22  
Expirydate: 2028.12.31



Following the procedure relating to the EC Declaration of Conformity set out in Annex II of  
Directive 93/42/EEC.

The above mentioned declaration of conformity is exclusively under the responsibility of  
Shenzhen Comen Medical Instruments Co.,Ltd

Company: Shenzhen Comen Medical Instruments Co.,Ltd

Address: Floor 10, Floor 11 and Section C of Floor 12 of Building 1A & Floor 1  
to Floor 5 of Building 2, FIYTA Timepiece Building, Nanhuan Avenue,  
Matian Sub-district, Guangming District, Shenzhen, Guangdong,  
518106, P.R. China.

2023.6.14  
Place, date

Legally binding signature, Function

Nr.	Basic UDI-DI	Denumire generică (denumirea dispozitivului)	Denumire comercială (brand)*	Modelul	Cod GMDN*
1	69454290DM001K6	Defibrilator Monitor	Comen	S5	17882



Administrator: Cobzari Turcan Rodica





# SHENZHEN COMEN MEDICAL INSTRUMENTS CO.,LTD

No.2 of FIYTA Timepiece Building, Nanhuan Avenue, Gongming  
Sub-district,Guangming New District, Shenzhen, P.R.China  
Web: www.comen.com E-mail: info@szcomen.com  
Tel: +86-755-26408879 Fax: +86-755-26431232

We, Shenzhen Comen Medical Instruments CO., LTD, having factories at No.2 of FIYTA Timepiece Building, Nanhuan Avenue, Gongming Sub-district, Guangming New District, Shenzhen, P.R.China, assign Aelo Grup SRL, based in Petru Movila str, 23/5, of.3 Chisinau MD -2004, Republic of Moldova, as **authorized representative** in correspondence with the conditions if directive 93/42/EEC, 98/79/EEC and 90/385/EEC.

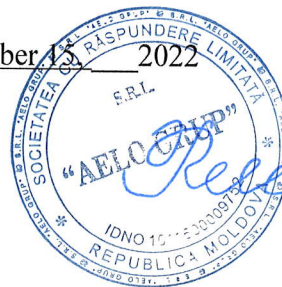
We declare that the company mentioned above is authorized to participate in public procurements with our medical equipment, to register, notify, renew or modify the registration of our medical devices on the territory of the Republic of Moldova.

Place: Shenzhen, China

Date: December 12 2022

Signed: \_\_\_\_\_

胡奇升



**EC Certificate**  
**Directive 93/42/EEC Annex II, excluding Section 4**  
**Full Quality Assurance System**  
**Medical Devices**

**Registration No.:** HD 60150327 0001

**Report No.:** 17058047 008

**Manufacturer:** SHENZHEN COMEN MEDICAL  
INSTRUMENTS CO., LTD.  
F10-11& Sect C, F12 of BLDG 1A and  
F1-5 of BLDG 2, FIYTA Timepiece  
Building, Nanhuan Avenue, Matian  
Subdistrict, Guangming District  
518106 Shenzhen, Guangdong  
P.R. China

**Products:** Medical Devices

(see attachment for products included)

Replaces Approval, Registration No.: HD 60144776 0001


**Expiry Date:** 2024-05-26

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

**Effective Date:** 2020-08-03

**Date:** 2020-08-03

Notified Body

  
Dipl.-Ing. W. Hsu



**TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg**  
TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC  
concerning medical devices with the identification number 0197.

**TÜV Rheinland**  
**LGA Products GmbH**  
**Tillystraße 2, 90431 Nürnberg**

Doc. 1/1 Rev. 0

**Attachment to  
Certificate**

**Registration No.:** HD 60150327 0001  
**Report No.:** 17058047 008

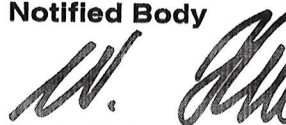
**Manufacturer:** SHENZHEN COMEN MEDICAL  
INSTRUMENTS CO., LTD.  
F10-11& Sect C, F12 of BLDG 1A and  
F1-5 of BLDG 2, FIYTA Timepiece  
Building, Nanhuan Avenue, Matian  
Subdistrict, Guangming District  
518106 Shenzhen, Guangdong  
P.R. China

**Products:**

- Anaesthetic Systems
- Syringe Pumps
- Infusion Workstation
- Infusion Pumps
- Neonatal Ventilators
- Medical Oxygen-air Blenders
- Infant Incubators
- Defibrillator/Monitors

**Date:** 2020-08-03

**Notified Body**

  
**Dipl.-Ing. W. Hsu**





# SGS

Certificate CN15/30544  
The management system of

## Shenzhen Comen Medical Instruments Co., Ltd.

Business Registration Address: Floor 10, Floor 11 and Section C of Floor 12 of Building 1A  
& Floor 1 to Floor 5 of Building 2, FIYTA Timepiece Building, Nanhuan Avenue, Matian  
Sub-district, Guangming District, Shenzhen, 518106, Guangdong, P.R. China  
Business Operation Address: Floor 10, Floor 11 and Section C of Floor 12 of Building 1A  
& Floor 1 to Floor 5 of Building 2, FIYTA Timepiece Building, Nanhuan Avenue, Matian  
Sub-district, Guangming District, Shenzhen, 518106, Guangdong, P.R. China

Unified Social Credit Code 91440300738806174Y  
has been assessed and certified as meeting the requirements of

### ISO 9001:2015

For the following activities

Design, Production and Sales of Fetal & Maternal Monitor, Multi-parameter Patient Monitor,  
Specialized Cardiovascular Monitor, Vital Signs Monitor, Specialized Neonatal Monitor,  
Specialized Fetal & Maternal Monitor, Defibrillator Monitor, Central Monitoring System  
Software, Anaesthetic Gas Scavenging System, Infant Incubator, T-piece Infant Resuscitation  
System, Electrocardiograph, Infrared Ear Thermometer, Anaesthesia Machine, Infusion Pump,  
Infusion Workstation, Syringe Pump, Neonatal Ventilator, Medical Oxygen-air Blender, Medical  
Air Compressor, Ceiling Pendant, LED Surgical Light, Infant Radiant Warmer, Infant  
Phototherapy Equipment, Catheter-positioning Guiding System, Temperature Control System,  
Video Laryngoscope, Sequential Compression System, High Flow Heated Respiratory  
Humidifier, Emergency and Transport Ventilator, Ventilator  
Design and Sales of Sterile disposable laryngoscope blade

This certificate is valid from 30 April 2021 until 29 April 2024  
and remains valid subject to satisfactory surveillance audits.

Recertification audit due a minimum of 60 days before the expiration date.

Issue 7. Certified since 30 April 2015

Authorised by

SGS United Kingdom Ltd  
Rossmore Business Park, Ellesmere Port, Cheshire, CH65 3EN, UK  
t +44 (0)151 350-6666 f +44 (0)151 350-6800 [www.sgs.com](http://www.sgs.com)

The certification information can be verified on the web site of Certification and Accreditation  
Administration of the People's Republic of China [www.cnca.gov.cn](http://www.cnca.gov.cn)



HC SGS 9001 2015 0118  
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of the law.





**Certificate CN15/30545**

### The management system of

# Shenzhen Comen Medical Instruments Co., Ltd.

Floor 10, Floor 11 and Section C of Floor 12 of Building 1A & Floor 1 to Floor 5 of Building 2, FIYTA Timepiece Building, Nanhuan Avenue, Matian Sub-district, Guangming District, Shenzhen, Guangdong, 518106, P.R. China

has been assessed and certified as meeting the requirements of



**ISO 13485:2016**  
**EN ISO 13485:2016**

**For the following activities**

**The scope of registration appears on page 2 of this certificate.**

**This certificate is valid from 02 March 2021 until 29 April 2024  
and remains valid subject to satisfactory surveillance audits.**

Re certification audit due before 04 February 2024  
Issue 8. Certified since 30 April 2015

Authorised by



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**SGS United Kingdom Ltd**  
**Rossmore Business Park Ellesmere Port Cheshire CH65 3EN UK**  
**t +44 (0)151 350-6666 f +44 (0)151 350-6600 [www.sgs.com](http://www.sgs.com)**

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## Shenzhen Comen Medical Instruments Co., Ltd.

**ISO 13485:2016**  
**EN ISO 13485:2016**



Issue 8

Detailed scope

### Design, Manufacture and Distribution of

- Electrocardiograph,
- Fetal & Maternal Monitor,
- Multi-parameter Patient Monitor,
- Specialized Cardiovascular Monitor,
- Vital Signs Monitor,
- Specialized Neonatal Monitor,
- Specialized Fetal & Maternal Monitor,
- Central Monitoring System Software
- Infrared Ear thermometer
- Anaesthetic Gas Scavenging System
- LED Surgical Light
- Ceiling Pendant
- T piece Infant Resuscitation System
- Anesthesia Machine,
- Infant Radiant Warmer.
- Infant Phototherapy equipment
- Catheter-positioning guiding system (Including Sterile Disposable electrode with extension wire)
- Syringe Pump
- Temperature Control System for management patient body temperature and vital physiological parameters Monitor
- Video Laryngoscope
- Sterile Disposable Laryngoscope blade
- Sequential Compression System for Prophylaxis of Vein Thrombosis and for alleviation of limb venous edema and pain
- Defibrillator Monitor
- High flow heated respiratory humidifier
- Emergency and Transport Ventilator
- Ventilator
- Operating table



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